

The Borgs, a New Family of Cdc42 and TC10 GTPase-Interacting Proteins

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Received 21 April 1999/Returned for modification 9 June 1999/Accepted 30 June 1999

The Rho family of GTPases plays key roles in the regulation of cell motility and morphogenesis. They also regulate protein kinase cascades, gene expression, and cell cycle progression. This multiplicity of roles requires that the Rho GTPases interact with a wide variety of downstream effector proteins. An understanding of their functions at a molecular level therefore requires the identification of the entire set of such effectors. Towards this end, we performed a two-hybrid screen using the TC10 GTPase as bait and identified a family of putative effector proteins related to MSE55, a murine stromal and epithelial cell protein of 55 kDa. We have named this family the Borg (binder of Rho GTPases) proteins. Complete open reading frames have been obtained for Borg1 through Borg3. We renamed MSE55 as Borg5. Borg1, Borg2, Borg4, and Borg5 bind both TC10 and Cdc42 in a GTP-dependent manner. Surprisingly, Borg3 bound only to Cdc42. An intact CRIB (Cdc42, Rac interactive binding) domain was required for binding. No interaction of the Borgs with Rac1 or RhoA was detectable. Three-hemagglutinin epitope (HA₃)-tagged Borg3 protein was mostly cytosolic when expressed ectopically in NIH 3T3 cells, with some accumulation in membrane ruffles. The phenotype induced by Borg3 was reminiscent of that caused by an inhibition of Rho function and was reversed by overexpression of Rho. Surprisingly, it was independent of the ability to bind Cdc42. Borg3 also inhibited Jun kinase activity by a mechanism that was independent of Cdc42 binding. HA₃-Borg3 expression caused substantial delays in the spreading of cells on fibronectin surfaces after replating, and the spread cells lacked stress fibers. We propose that the Borg proteins function as negative regulators of Rho GTPase signaling.

Motility and morphogenesis are probably among the most complicated processes that a cell performs. The proteins and molecular mechanisms that regulate these processes are only now beginning to be elucidated. Adherent cells elaborate an extracellular matrix of proteins to which they bind through receptors called integrins (8). The integrins cluster at focal adhesion complexes and transmit signals through these complexes to the actin-myosin cytoskeleton (11; for reviews, see references 10 and 46). Signaling through other types of receptor, such as those that bind growth factors, can be modulated by engagement of integrins with the extracellular matrix (39). A key role in controlling focal adhesions and the actin cytoskeleton is played by the small, Rho-like GTPases (9, 19, 28, 29), of which there are at least 13 types in mammalian cells (15, 36, 54). The most intensively studied members of this family of GTPases are RhoA, Rac1A, and Cdc42. Many of these proteins induce dramatic changes in the actin cytoskeleton when expressed ectopically as gain-of-function mutants (23, 35, 40, 41; reviewed in reference 16) and can perturb cell adhesion, cell spreading, motility, and cytokinesis (11, 20, 44, 50; reviewed in reference 26 and 55). The replating of fibroblasts from suspension culture onto a fibronectin-coated surface causes dramatic membrane ruffling and the rapid production of lamellipodia and microspikes around the edges of the spreading cells. These changes are mediated by the activation of the Rac and Cdc42 GTPases (5). The RhoA GTPase is transiently inhibited after replating and then activated at a later stage of spreading, at which time actin stress fibers appear within the cytosol (11). Similar changes in the activities of these

proteins may occur at the leading edge of motile cells. Additionally, the Rho GTPases can activate protein kinase cascades and transcription factors and can regulate entry into the cell cycle (2, 12, 24, 37, 55).

Given this wealth of responses, it is not surprising that each of the Rho family GTPases has been found to interact with a plethora of target proteins that likely act as downstream effectors. These proteins include a variety of types of kinase and of adapters, plus other proteins of unknown function (for reviews, see reference 51 and 55). Several of the adapter proteins interact with known components of the actin cytoskeleton such as profilin, but their roles remain unclear (49, 56).

Because the Rho family GTPases mediates changes in gene expression and cell division that are independent of the actin cytoskeleton, different subsets of effectors likely participate in distinct signal transduction pathways downstream of the GTPases (24, 52). It is only through the identification and detailed analysis of the complete set of Rho family GTPases and of their effector proteins, therefore, that we will achieve satisfactory understanding of the molecular basis for those aspects of motility and morphogenesis that are influenced by these GTPases.

Toward this end, we performed a large two-hybrid screen of a whole mouse embryo library, using the TC10 GTPase as bait. Although its cDNA was cloned almost a decade ago (14), TC10 has been characterized only recently (14, 33). Of the >250 positives clones that we isolated in the screen, many contained open reading frame fragments of previously described proteins that interact with Cdc42 (33). These proteins include the protein kinases α -PAK, β -PAK, γ -PAK, PAK4, myotonic dystrophy kinase-related Cdc42-binding kinase (MRCK)- α and - β , and mixed-lineage kinase 2 (MLK2) (1, 25, 48, 51). We also isolated a partial cDNA for a putative new MRCK protein that we called MRCK- γ . In addition, the screen identified N-WASP

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(30) and an uncharacterized protein called MSE55 (marrow stromal/endothelial cell protein with a molecular mass of 55,000 Da) (3). All of these clones contained a CRIB (Cdc42, Rac interactive binding) domain, which has been shown to be necessary and sufficient to permit association with the Cdc42 and/or Rac proteins when in the GTP-bound state (7). Three other downstream effectors of Cdc42 that also contain CRIB domains, namely, WASP, p120ACK, and MLK3 (51), were not identified in the two-hybrid screen. These proteins do not bind TC10 (33).

We now describe a new family of five related TC10 and Cdc42 binding proteins, which we have called Borgs (binders of Rho GTPases). Three of the cDNAs for these proteins were identified in the two-hybrid screen, and two more were uncovered by database searches. One protein of this family (Borg5) is identical to the previously reported protein MSE55 (3), which was discovered serendipitously as a protein cross-reacting with an anti-hemonection antibody. A Northern blot detected MSE55 mRNA only in endothelial and bone marrow stromal cells (3). The cDNA encodes a protein of 391 amino acid residues with two regions of weak similarity to some calcium binding domains. More recently, a database search revealed that the protein contains a CRIB domain, and purified glutathione *S*-transferase (GST)–MSE55 protein was shown to bind efficiently to [γ - 32 P]GTP–Cdc42 and more weakly to [γ - 32 P]GTP–Rac (7). We demonstrate that the other Borg family members are more widely expressed, that bind TC10 and/or Cdc42, and can inhibit the Cdc42-mediated activation of the c-Jun N-terminal kinase (JNK). Ectopic expression of Borg1 or Borg3 induces distinctive changes in cell morphology and significantly delays cell spreading on fibronectin as well as cell motility in a wound test.

MATERIALS AND METHODS

Cloning of Borg cDNAs. The two-hybrid screen has been described previously (33). The bait was TC10(Q75L) Δ C, a gain-of-function mutant of TC10 lacking the C-terminal isoprenylation signal and fused to the DNA binding domain of the yeast GAL4 protein. The library was from a mouse embryo (day 9) fused to the activating domain of VP16 and was kindly provided by Stan Hollenberg (Seattle, Wash.). Four positive clones encoded a protein sequence highly similar to that of the human protein MSE55 (amino acid residues 1 to 110) (3). Differences in 11 amino acid residues between the clones and MSE55 most likely represent interspecies differences.

A set of two identical clones and another of five clones each encoded new CRIB domains which showed closer similarity to that of MSE55 than to CRIB domains of other known Rho GTPase binding proteins. The complete human cDNA of the first gene was obtained from expressed sequence tag (EST) plasmid R13749/26651 and was named Borg1. Both fragments found in the screen correspond to amino acids 1 to 181 of the human open reading frame of Borg1 and encompassed a complete CRIB domain. Alignments of sequences were by made using the clustal algorithm in the Megalign program from the DNASTar software package. The mouse sequence from the screen possesses four additional amino acid residues (PSPO) between positions 143 and 144. Overall, the mouse and human sequences are 81% identical. The set of five clones encode a reading frame, which we have termed Borg4, for which no full-length ESTs were present in the database. A partial human sequence of Borg4 was obtained from EST AA013011/360199. The sequence encodes two additional amino acid residues (SS) located between positions 68 and 69 of the mouse sequence.

When the databases were searched by BLAST, using a region of the CRIB domains of Borg1 and MSE55/Borg5 as the query, we identified two new open reading frames for closely related proteins, which we named Borg2 and Borg3. EST AA046144/376758 encoded the complete human Borg2 protein, and EST W85265/408069 encoded the complete mouse Borg3 protein. In all cases, we have defined "complete" as meaning that the open reading frames begin with an initiation codon downstream of a Kozak consensus sequence (22) and that in-frame stop codons are found upstream of this sequence. Full-length Borg1, Borg2, and Borg3 cDNAs were mutagenized by PCR so as to create a *Bam*HI (or, for Borg2, a *Bgl*II) site at the 5' end and an *Eco*RI site at the 3' end of the open reading frames. These DNA fragments were then cloned in frame into the pKH3 (6), pGEX2T, and VP16 vectors that had been cut with *Bam*HI and *Eco*RI. The vectors produce fusion proteins with an N-terminal triple-HA1 (hemagglutinin epitope) tag, GST, or VP16 activation domain, respectively.

Two conserved residues within the CRIB domain of Borg3 (Ile²³ and Ser²⁴)

were mutated to Ala residues by megaprimer PCR mutagenesis (4), to produce CRIB (I₂₃A,S₂₄A). A truncation mutant in which the C-terminal 67 amino acid residues of Borg3 were deleted was produced by PCR. Recombinant proteins were expressed and purified from *Escherichia coli* as previously described (6).

Yeast conjugation assay. Conjugation assays were performed as previously described (33). Briefly, *Saccharomyces cerevisiae* HF7c (*MATa*) was transformed with VP16 vectors that express fragments or full-length sequences of Borg proteins fused to the VP16 activation domain. Strain W303 (*MATa*) was transformed with modified pGBT9 vectors expressing GAL4 DNA binding domain fusions with different activated mutants of Rho family GTPases lacking the C-terminal isoprenylation signal. The two mating types were mixed and incubated overnight on YPD plates to permit mating. The following day, the spots were replica plated onto selective medium (L⁻/W⁻/H⁻ plus 10 mM 3-aminotriazole) and incubated at 30°C to permit growth of diploids containing interacting proteins. Other known target proteins that interact with Rac(G12V) Δ C and Rho(G14V) Δ C were tested in the same assay as positive controls for expression of these GTPases.

In vitro binding assays. Recombinant GST–Borg proteins were purified, and 2 μ g of each, or of GST alone, was spotted onto nitrocellulose and allowed to dry for 1 h at room temperature. The nitrocellulose was blocked in binding buffer (50 mM Tris–HCl [pH 7.5], 100 mM NaCl, 5 mM MgCl₂, 0.1 mM dithiothreitol) plus 5% dried milk for 1 h at 4°C. TC10 and Cdc42 proteins (3 μ g) were loaded with [α - 32 P]GTP or [α - 32 P]GDP (3,000 Ci/mmol) as described previously (33) and separated from unincorporated nucleotide by passage over a Pharmacia PD10 size exclusion column equilibrated in binding buffer. Equal amounts of radiolabeled proteins (ca. 0.5 μ Ci of each) were incubated with the nitrocellulose in 10 ml of binding buffer for 1 h at 4°C. The blots were subsequently washed three times with 50 ml of binding buffer, and associated proteins were detected by fluorography.

Immunodetection, immunofluorescence, and imaging. NIH 3T3 fibroblasts were cultured on 10-cm-diameter plates or on two-well LabTek chamber slides (Nunc) in Dulbecco modified Eagle medium (DMEM) supplemented with 5% fetal calf serum, 5% calf serum, penicillin, and streptomycin. The cells were transfected by the calcium phosphate precipitation technique as previously described (43). For each transfection experiment, 12 μ g of DNA was mixed with the calcium phosphate reagents in a final volume of 1.2 ml. One-sixth of the mix was added per LabTek chamber (for immunofluorescence studies), and the remainder was added per 10-cm-diameter plate of cells (for immunoblotting). When two plasmids (12 μ g of each) are mixed and cotransfected, we find that under these conditions about 90% of the transfected cells express both proteins.

Cell protein extracts were prepared from the plates 48 h posttransfection, submitted to sodium dodecyl sulfate–polyacrylamide gel electrophoresis (SDS–PAGE) and transferred to nitrocellulose membranes as described previously. Proteins were detected with monoclonal antibody 12CA5 (anti-HA; 1/5,000) or 9E10 (anti-Myc; 1/20,000) and revealed with a horseradish peroxidase-conjugated anti-mouse secondary antibody by using chemiluminescence (Kirkegaard & Perry Laboratories, Gaithersburg, Md.). For immunofluorescence, cells were washed in phosphate-buffered saline (PBS) 40 h posttransfection and then fixed in paraformaldehyde (4% [wt/vol] in PBS) for 15 min at room temperature. Cells were washed in PBS, and residual formaldehyde was quenched with 50 mM ammonium chloride for 15 min. Cells were permeabilized with Triton X-100 (0.2% [vol/vol] in PBS) for 3 min and blocked in PBS plus 3% (wt/vol) bovine serum albumin for at least 20 min. Cells were then incubated with mouse monoclonal anti-HA antibody 12CA5 (1/500) in PBS plus 0.3% bovine serum albumin for 1 h. After being washed in the same buffer, cells were incubated for 45 min with the secondary, Texas red-coupled antibody (1/1,200) and fluorescein isothiocyanate (FITC)-phalloidin (1/1,000; Sigma). Slides were mounted in Citifluor (Ted Pella, Redding, Calif.) and imaged by using a 60 \times water immersion objective lens on a Nikon inverted microscope equipped with a Hamamatsu charge-coupled device camera. Data were captured and processed by using Openlab 2.0 (Improvision) and Adobe Photoshop 5.0 software.

JNK assay. Cos or NIH 3T3 cells, grown in DMEM supplemented with 5% fetal calf serum, 5% calf serum, penicillin, and streptomycin, were cotransfected with plasmids as described above. Control vectors pKH3 and pRK7 were used as necessary to normalize the amount of DNA transfected to 9 μ g per 100-mm-diameter plate. Cells were incubated overnight in serum-free DMEM 24 h posttransfection; then 100 μ g of anisomycin was added (control), and cells were incubated for 20 min at 37°C. Cells were washed twice with 10 ml of ice-cold PBS and lysed in 400 μ l of lysis buffer (25 mM HEPES [pH 7.4], 300 mM NaCl, 1.5 mM MgCl₂, 0.5 mM dithiothreitol, 20 mM β -glycerophosphate, 1 mM Na₃VO₄, 0.1% Triton X-100, 20 μ g of aprotinin/ml, 10 μ g of leupeptin/ml, 1 mM phenylmethylsulfonyl fluoride, 1 μ M okadaic acid). Lysed cells were scraped off the plates, and the supernatant was cleared by centrifugation (5 min at 14,000 \times g, 4°C). Equal volumes (100 μ l) of the supernatant were removed for immunoblot analysis and separated on 12% polyacrylamide gels by SDS–PAGE. The proteins were transferred to a nitrocellulose membrane and immunoblotted. HA₃-tagged proteins were immunoprecipitated from the remaining supernatant and washed three times in buffer A (2 mM Na₃VO₄ and 1% NP-40 in PBS), once in buffer B (100 mM morpholinepropanesulfonic acid [MOPS; pH 7.5], 0.5 M LiCl), and once in kinase buffer (12.5 mM MOPS [pH 7.5], 12.5 mM β -glycerophosphate, 7.5 mM MgCl₂, 0.5 mM EGTA, 0.5 mM NaF, 0.5 mM Na₃VO₄). Immunoprecipitated proteins were resuspended in 30 μ l of kinase buffer, and 2 μ g of purified

GST—c-Jun(1–79) plus 2 μ Ci of [γ - 32 P]ATP were added to each reaction tube. The reactions proceeded at 30°C for 20 min and were stopped by the addition of 10 μ l of 4 \times Laemmli sample buffer. Kinase reaction complexes were separated on 12% polyacrylamide gels by SDS-PAGE and analyzed by fluorography.

Cell spreading and motility assays. NIH 3T3 cells in 10-cm-diameter plates were transfected with 10 μ g of DNA. Forty hours after transfection, cells were trypsinized, maintained in suspension for 15 to 20 min, and then replated on LabTek chambers that had been coated with fibronectin (0.5 ml of a 50- μ g/ml solution). At different times (18, 45, or 180 min) after replating, cells were fixed with paraformaldehyde and processed for immunofluorescence as described above. Transfected cells were analyzed by using Openlab 2.0 software to measure cell surface areas in contact with the fibronectin surface. For each time point, the areas of 100 cells were counted. To compare expression levels of the Borg proteins within the transfected cells, images were captured under conditions such that no camera pixels were saturated, and the mean fluorescence of the counted cells was determined for each experiment. Differences in mean cell area at 18 min postplating were analyzed by an unpaired *t* test assuming equal variances (variances were calculated for each data set and found to be approximately equal). Probabilities (*P* values) were also calculated for the *t*-test values and degrees of freedom (i.e., 198).

To quantitate cell motility, a wounding assay was used. Cells were cotransfected with the plasmid of choice plus pK7-GFP, which expresses a bright mutant of green fluorescent protein (GFP). Cells were split and replated at high density and low density onto two 35-mm-diameter plates. After 3 days, when the cells were confluent, an aspirator was drawn across the center of the high-density culture, to produce a clean 1-mm-wide wound area free of cell debris. After a further 24 h of incubation to permit migration of cells into the empty area of the plate, the numbers of green and nonfluorescent cells in the cleared area were counted and compared to the ratio of green to nonfluorescent cells in the low-density, unwounded culture. Cells cotransfected with pK7-GFP and the pRK7 empty plasmid gave a ratio close to 1.0, a value expected if the GFP itself has no effect on motility. Statistical significance of the data was assessed by a two-sample *t* test.

Nucleotide sequence accession numbers. Accession numbers for the Borg open reading frame sequences are AF163840 (Borg1), AF164118 (Borg2), AF164119 (Borg3), and AF165114 (Borg4).

RESULTS

Cloning of Borg family cDNAs. A two-hybrid screen of a 9-day mouse embryo library using an activated TC10 GTPase as bait identified numerous clones that encoded protein fragments incorporating a CRIB domain. Three groups of clones had almost identical CRIB domains and similar flanking sequences, suggesting that they belonged to a family of related proteins that interact with Rho family GTPases (Fig. 1). Database searches using the protein sequence of each group uncovered five sets of human ESTs encoding proteins with a high degree of similarity to the two hybrid clones. One of these clones was almost identical to the previously reported human MSE55 protein (3) and is most likely the mouse counterpart of MSE55. We obtained from different ESTs the complete open reading frames for three proteins of this new family but found no ESTs encompassing the 3' end of one cDNA found in the screen.

Because the predicted sizes of the members of this family are all lower than 55 kDa, because the tissue expression of the corresponding mRNAs is not restricted to endothelial and stromal cells (see below), and because the name MSE55 does not indicate function, we decided to rename the whole family Borg proteins. Borg1 is the complete new protein of which a part was revealed in the two-hybrid screen, Borg2 and Borg3 are the two proteins for which DNA sequences were found in the EST database, Borg4 corresponds to the incomplete cDNA identified in the screen, and Borg5 is MSE55. Two uncharacterized sequences which correspond to Borg1 have been deposited by others in the GenBank database (accession no. AF001436 and AF094521).

Surprisingly, the mouse sequence of Borg1 contains four additional amino acids after position 143 that are a repeat of the four previous amino acids (PSPQ). This discrepancy is confirmed in two other independent mouse EST clones. Additionally, the human sequence of Borg4 contains an insertion

of two amino acids between positions 68 and 69 of the mouse sequence (SSSK versus SK). These differences in otherwise well-conserved sequences might reflect the presence of different splice variants of the same proteins.

The domain structures of the Borg proteins are diagrammed in Fig. 1A. Borg1, Borg2, and especially Borg3 (210 [22.5 kDa], 254 [27.5 kDa], and 150 [15.5 kDa] amino acid residues, respectively) are smaller than Borg5/MSE55 (which consists of 379 amino acid residues, for a predicted molecular mass of 39 kDa). Bahou et al. (3) described MSE55 as a 391-amino-acid protein, but the initiation codon that they indicated corresponds, at the mRNA level, to a very unlikely translation start. The Kozak context (22) for translation is unfavorable: CUGA UGC versus the best consensus described, ACCAUGG (the first adenine being present in 90% of cases) (initiation codon indicated by underlining). The context of the second in-frame Met codon (ACCAUGA) is closer to the Kozak consensus motif and aligns naturally with the initiation codon of the other Borg proteins (Fig. 1B). We suggest, therefore, that it most probably represents the true start site for translation of Borg5 and have numbered residues in Fig. 1 accordingly. The N-terminal regions of all five Borg proteins are highly similar, comprising a short basic region (from 13 amino acids for Borg1 to as short as 3 amino acids for Borg5) followed by a CRIB domain sequence, ISXPLGDFRHTXH(I/V)G. This sequence matches the consensus motif, I(S/G)XPXXFXHXXHVG (7), with an additional residue in the Borg CRIB domains between the P and F residues.

A well-conserved, short domain of about 12 amino acids that is not found in any other known protein follows the CRIB domain. We therefore call it Borg homology 1 (BH1) domain. The central and C-terminal parts of the proteins are more divergent. Two other well-conserved motifs that we call the BH2 and BH3 domains can be defined (Fig. 1). The BH2 domain is not present in Borg3; the BH3 domain has a central location in Borg5, whereas it is localized at the C-terminal parts of Borg1, Borg2, and Borg3. Neither domain is present within the N-terminal fragment of Borg4 for which we have sequence information. A proline-rich domain (about 40% of Pro in each case) can be observed in the central regions of Borg1, Borg3, and Borg5. The four amino acid residues insertion of the mouse Borg1 sequence is located within this region. Borg5 also contain eight in-tandem heptad repeats (consensus PAANPPA) located at the C-terminal side of the BH3 domain.

Borg mRNAs show distinct expression patterns. The MSE55 gene is expressed in endothelial and bone marrow stromal cells but not in monkey liver, spleen, brain, lung, and kidney (3). To determine the tissue expression patterns of other members of the family, fragments of cDNA encoding human Borg1, Borg2, or Borg4 or mouse Borg3 were used to probe a human tissue poly(A)⁺ mRNA blot (Fig. 2). *Borg1*, *Borg2*, and *Borg3* mRNAs were detected in all eight tissues studied, though at very different levels. For all three genes, multiple transcripts were observed. *Borg1* is expressed as two transcripts of about 1.8 and 2 kb and at high levels in heart. *Borg1* mRNA is less abundant in pancreas and especially liver. The Borg2 probe revealed four different transcripts of various sizes (from 2 to 5.5 kb). *Borg2* is also expressed at high levels in heart but at low levels in brain, and the largest transcript was not detected in liver. The Borg3 probe was not of human origin and gave a weaker signal. However, transcripts of about 1 and 2 kb are visible (Fig. 2) and were more clearly seen in a rat cardiomyocyte mRNA extract (data not shown). In addition to these two messengers, present in all examined tissues, a more abundant transcript of about 3.5 kb is detectable specifically in skeletal muscle. The presence of multiple transcripts suggests the pos-

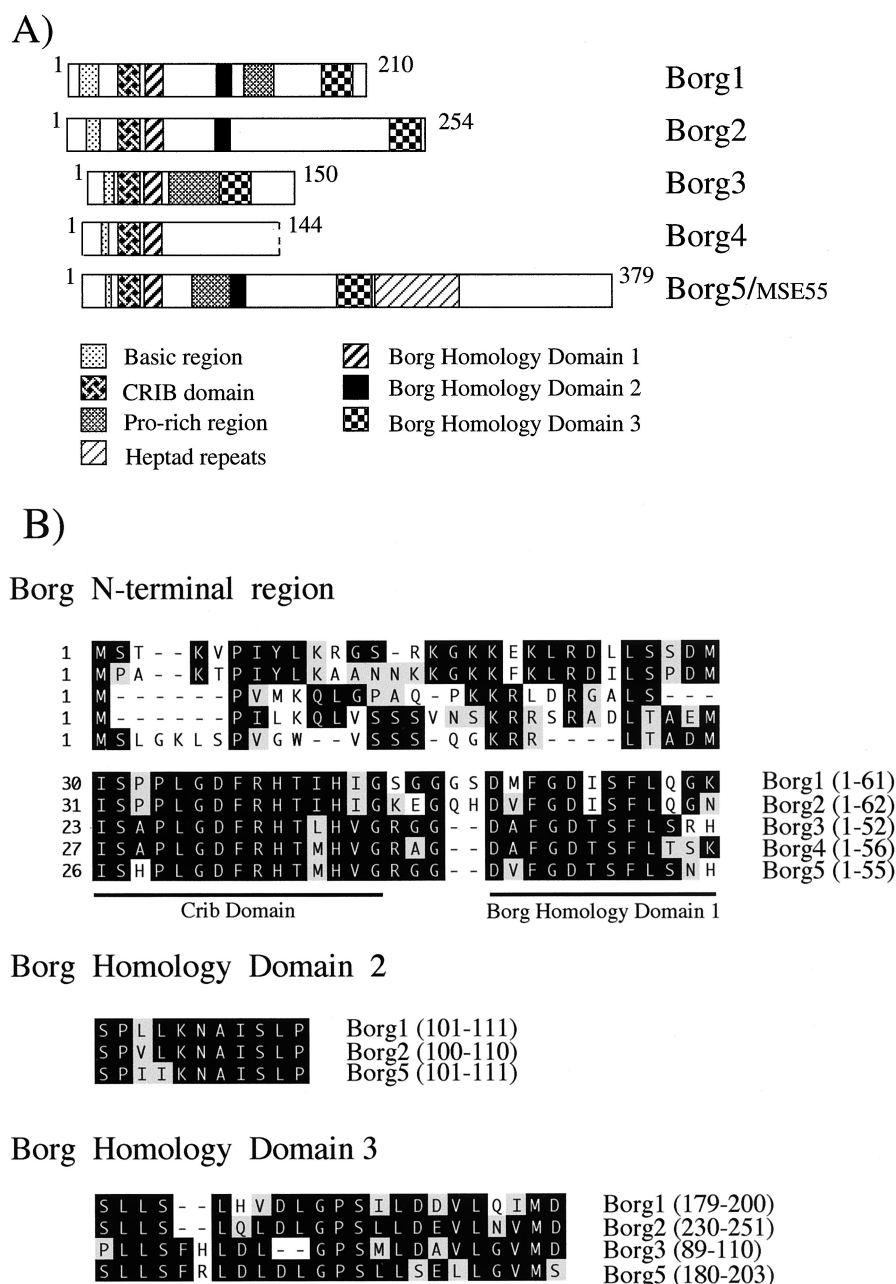


FIG. 1. (A) Schematic of Borg protein domain structures. Bars showing the regions of similarity within the primary sequences of the Borg proteins are aligned according to the CRIB domains. The C-terminal sequence of Borg4 is not known. Borg5 is the putative mouse homolog of human protein MSE55. (B) Alignment of BH1, BH2, and BH3 domains. The N-terminal region comprises the CRIB domain and the BH1 domain.

sible existence of differentially spliced mRNAs and of Borg protein variants. Indeed, two slightly different mRNAs of *MSE55* have been described (3). No transcripts were found in any tissues probed with the Borg4 probe. The *Borg4* gene might, like *Borg5*, be expressed in only a few, specific tissues or cell types. However, it is striking that all available ESTs are of embryonic or fetal origin. Therefore, Borg4 may not be expressed in adult tissues.

Borg3 interacts specifically with Cdc42. Partial cDNAs for Borg1, Borg4, and Borg5 were found in the two-hybrid screen. All clones of these cDNAs contained the 5' coding region including the region corresponding to the CRIB and BH1

domains of the proteins. In a yeast two-hybrid conjugation assay, the expressed protein fragments of Borg1, Borg4, and Borg5 interacted specifically with the activated TC10 GTPase deleted of its isoprenylation signal [TC10(Q75L) Δ C] and not with the GDP-bound TC10 mutant [TC10(T31N) Δ C] (Fig. 3A shows data for Borg4 and Borg5). They also bound to an activated mutant of Cdc42 [Cdc42(Q61L) Δ C]. To check that the interaction was not an artifact of the use of fragments rather than full-length proteins, we repeated the conjugation assay using the complete open reading frames of Borg1 to Borg3. Surprisingly, although the Borg1 and Borg2 interacted specifically with TC10, Borg3 did not (Fig. 3A). Borg3 did,

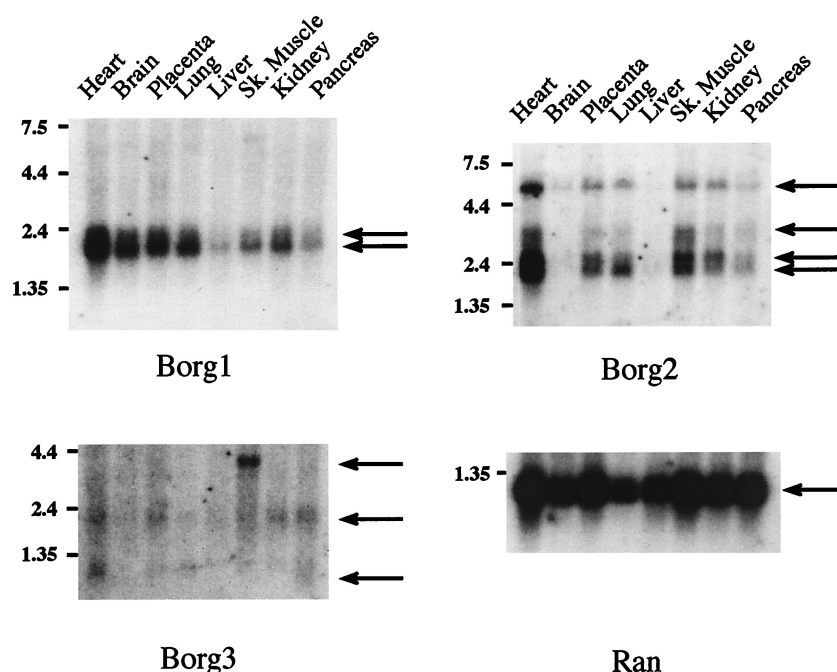


FIG. 2. Tissue-specific expression of *Borg* genes. A nylon membrane containing human poly(A)⁺ mRNAs from different tissues (Clontech) was hybridized with α -³²P-labeled human (mouse for Borg3) cDNA probes encoding the Borg proteins. Ran GTPase cDNA was used as a loading control. The same membrane was used in all four Northern blots. Molecular sizes of RNA standards are indicated (in kilobases) on the left. Arrows indicate the multiple transcripts that hybridized with each probe.

however, interact strongly with Cdc42. Burbelo et al. reported a weak binding of MSE55/Borg5 to GTP-loaded Rac1 in a dot blot assay (7), but in our hands, no member of the Borg family gave a positive response in the two-hybrid assay with activated mutants of either Rac1 or RhoA (Fig. 3A).

To determine whether the observed interactions between the small GTPases and the Borgs is direct, we spotted equal amounts of recombinant GST-Borg1, GST-Borg2, and GST-Borg3 fusion proteins onto nitrocellulose and incubated them with recombinant TC10 or Cdc42 that had been loaded with [α -³²P]GTP. After washing, the GST-Borg1 and GST-Borg2 proteins showed strong binding to both Cdc42 and TC10. Binding of GST-Borg3 to TC10 was undetectable, although it bound the GTP-loaded Cdc42 with high affinity (Fig. 3B). [α -³²P]GDP-Cdc42 did not bind to GST-Borg3, demonstrating that, as expected, the binding of Cdc42 to Borg3 is nucleotide dependent (Fig. 3C). These results confirm the validity of the yeast two-hybrid conjugation assay and demonstrate that the original two-hybrid screen was unable to identify Borg3 not because it was absent from the library but because it does not recognize TC10. It is interesting that the Borg3 CRIB domain, though closely related to the CRIB domains of the other Borg family members, nonetheless contains determinants that can differentiate TC10 from Cdc42, despite the fact that these GTPase effector domain sequences are almost identical. Similarly, we have found that while MLK2 and N-WASP interact with TC10, MLK3 and WASP do not, though all seem to bind with similar affinity to Cdc42 (33).

To confirm that the CRIB domain constitutes the only binding site on Borg proteins for Cdc42, we created a GST-Borg3 mutant in which the two first, highly conserved amino acid residues (Ile and Ser) of the CRIB motif were converted to Ala residues [Borg3(I₂₃A,S₂₄A)]. This fusion protein did not detectably bind [α -³²P]GTP-loaded Cdc42 (Fig. 3B). We also created a C-terminal truncation mutant of Borg3, GST-

Borg3(1–83), which lacks BH3 and part of the Pro-rich domain. This mutant bound Cdc42 as well as did the wild-type protein. These results argue that the CRIB domain region of Borg3 is necessary and sufficient for interaction with the Cdc42 GTPase.

Borg3 inhibits JNK independently of Cdc42 binding. JNK is the terminal protein kinase of a cascade that can be activated by numerous stress-related signal inputs and by a variety of seven-transmembrane domain receptors that couple to heterotrimeric G proteins (31). The $\beta\gamma$ subunits released from the G protein upon activation of these receptors are believed to activate exchange factors for Rho family GTPases (13). Cdc42 can then activate the protein kinase cascade that switches on JNK. The mixed-lineage kinase MLK2 may mediate this activation (18). However, there remains the possibility that other downstream effectors of Cdc42 participate in the pathway. We therefore tested the effects of Borg expression on the activity of JNK in transfected Cos cells.

As shown in Fig. 4, the coexpression of HA₃-Borg3 suppressed the basal activity of HA-JNK. Anisomycin stimulated both the basal JNK and Borg-inhibited JNK activities by about the same degree, indicating that Borg3 does not interfere with the stress-mediated signaling pathway that regulates JNK. The inhibition by HA₃-Borg3 was not a result of changes in the level of expression of the HA-JNK or of competition for the anti-HA antibody. In fact, HA₃-Borg3 is not efficiently immunoprecipitated by the anti-HA₃ tag antibody, perhaps because the tag is hidden by interaction of the Borg with other proteins or because it is partially buried within the folded protein (not shown). To assess whether this inhibition is dependent on the binding of Cdc42, we used in the same assay a CRIB domain mutant, Borg3(I₂₃A,S₂₄A), that cannot bind Cdc42. Expression of this mutant inhibited basal JNK activity to the same extent as the wild-type Borg3 (Fig. 4B). In each case, we checked the presence of all ectopically expressed proteins in the lysates by

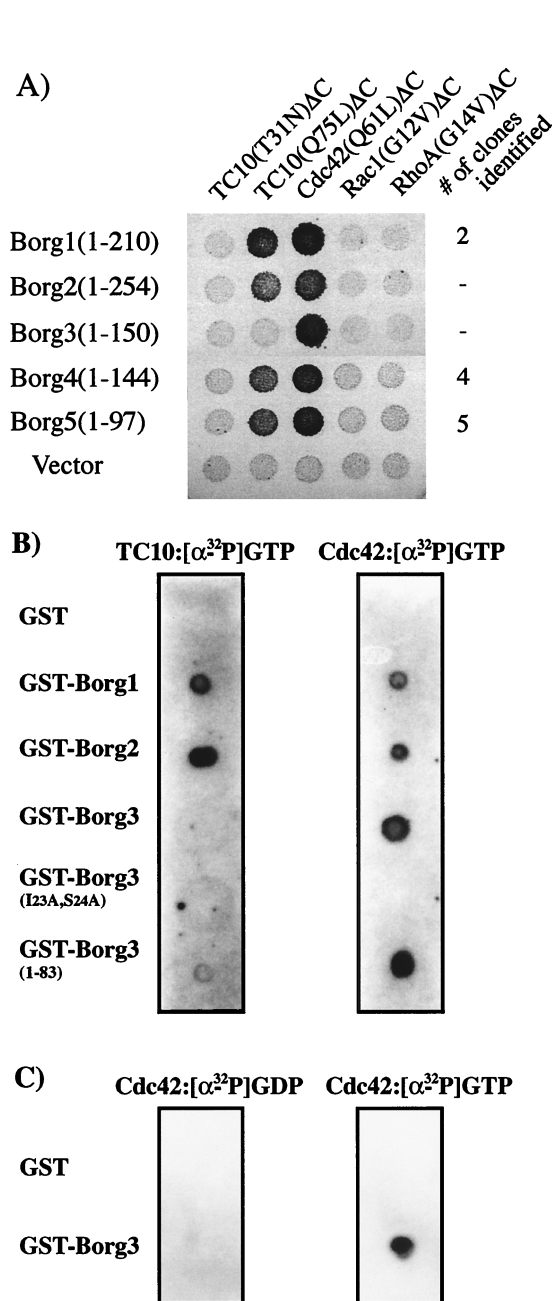


FIG. 3. Binding of Borg proteins to different Rho-like GTPases. (A) Yeast two-hybrid conjugation assays. *S. cerevisiae* HF7c (*MATa*) was transformed with VP16 empty vector or with VP16 containing Borg cDNAs (the complete open reading frame of Borg1, Borg2, or Borg3 or a fragment obtained from the two-hybrid screen for Borg4 and Borg5). The transformants were then conjugated to *S. cerevisiae* W303 (*MATα*) which had been transformed with pGBT9 plasmids containing cDNA encoding different Rho-like GTPases. Each GTPase open reading frame had been truncated to remove the C-terminal isoprenylation signal. Results show the growth of diploids after replica plating onto selective medium ($L^{-}/W^{-}/H^{-}$ plus 10 mM 3-aminotriazole). For the three Borg cDNAs identified in the screen, the number of independent clones identified is noted. (B and C) Binding of GST-Borg proteins to TC10-GTP, Cdc42-GTP, or Cdc42-GDP. Purified recombinant GST or GST-Borg1, GST-Borg2, GST-Borg3, GST-Borg3(I23A,S24A), and GST-Borg3(1-83) (2 μ g of each) were spotted onto a nitrocellulose membrane, which was then incubated with TC10-[α -³²P]GTP, Cdc42-[α -³²P]GTP, or Cdc42-[α -³²P]GDP (0.5 μ Ci; 3,000 Ci/mmol). Interactions were revealed by fluorography after the membranes were washed in binding buffer as described in Materials and Methods.

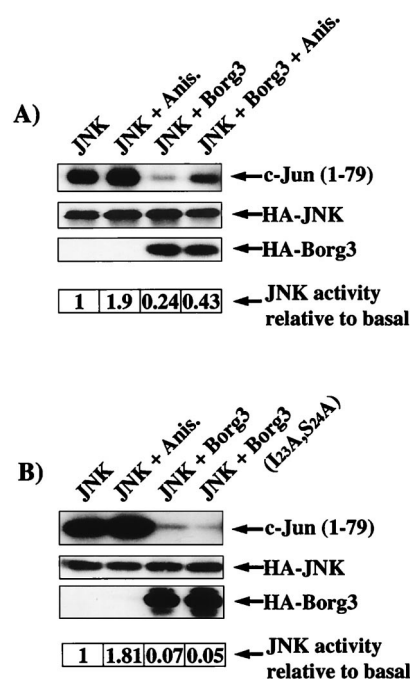


FIG. 4. Inhibition of JNK by Borg3. Cos cells were transfected as described in Materials and Methods and incubated overnight in serum-free DMEM. After 24 h, 100 μ g of anisomycin (Anis.) was added (control) and the cells were incubated for 20 min at 37°C. Cells were lysed in 400 μ l of lysis buffer. After clearing, equal volumes (100 μ l) of the supernatant were removed for immunoblot analysis and separated by SDS-PAGE. Proteins were transferred to a nitrocellulose membrane and immunoblotted as described in Materials and Methods. HA₃-tagged proteins were immunoprecipitated from the remaining supernatant and resuspended in 30 μ l of kinase buffer, and 2 μ g of purified GST-c-Jun(1-79) as the substrate plus 2 μ Ci of [γ -³²P] ATP were added to each reaction tube. The reactions proceeded at 30°C for 20 min and were stopped by the addition of 10 μ l of 4 \times Laemmli sample buffer. Kinase reaction complexes were separated by SDS-PAGE and analyzed by fluorography.

immunoblotting (Fig. 4). A similar inhibition of JNK activity by HA₃-Borg3 and HA₃-Borg3(I23A,S24A) was observed in transfected NIH 3T3 cells (not shown).

These data indicate that the inhibition of the JNK pathway by Borg proteins is independent of the binding to Rho-like GTPases and therefore is not a result of competition for another downstream effector such as MLK2, which is required for JNK activation.

Effects of ectopic expression of Borg proteins on cell shape. To examine whether Borgs might, like many downstream effectors of Cdc42/Rac, play a role in the elaboration or remodeling of the actin cytoskeleton, we expressed Borg1, Borg2, and Borg3 as HA₃-tagged fusion proteins in NIH 3T3 fibroblasts. As a negative control, cells were transfected with the empty vector, pKH₃. Expression of the Borg protein was detected with the monoclonal anti-HA antibody 12CA5 and a secondary Texas red-coupled antibody. The F-actin framework was revealed by using FITC-phalloidin. Expression of HA₃-Borg1 did not cause any dramatic changes in cell shape, but there was a substantially reduced abundance of stress fibers in most of the transfected cells, and unusually long, thin extensions were common (Fig. 5b and c). Similar but more pronounced morphological changes were seen when we ectopically expressed HA₃-Borg2 and especially HA₃-Borg3. Most of the transfected cells displayed a disturbed shape, in which the cell bodies had pulled in, but also extended long, sometimes beaded processes and protrusive lamellipodia (Fig. 5d to g). This phenotype is

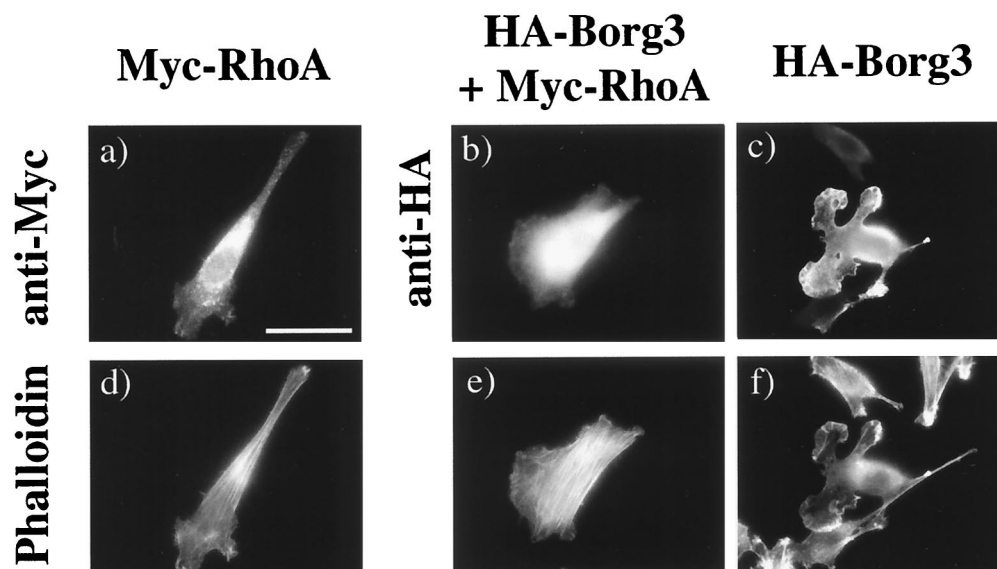


FIG. 6. Borg phenotype is abolished by coexpression of Myc-RhoA. NIH 3T3 fibroblasts were transfected with either pKMyC containing cDNA encoding for RhoA protein (a and d) or pKH₃ containing cDNA encoding Borg3 protein (c and f) or were cotransfected with plasmids encoding HA₃-Borg3 and Myc-RhoA proteins (b and e). Cells were fixed and processed for immunofluorescence. Tagged proteins were detected with either monoclonal anti-HA antibody 12CA5 (b and c) or monoclonal anti-Myc antibody 9E10 (a). Both antibodies were revealed with a secondary, Texas red-coupled anti-mouse antibody. F-actin was revealed with FITC-phalloidin (d to f). Bar = 40 μ m.

Myc-tagged RhoA alone, or with HA₃-Borg3 plus Myc-RhoA. As can be seen in Fig. 6c and f, the Borg3 produced a characteristic distortion of the normal cell morphology, with loss of stress fibers and the extension of protrusive lamellipodia. Coexpression of the Myc-RhoA dramatically increased stress fiber formation and the restoration of a more normal fibroblast appearance. This result supports the hypothesis that Borg3 inhibits Rho function and suggests that it acts upstream of or parallel to Rho rather than by inhibiting a downstream effector of Rho.

Finally, to determine whether the effect of the Borg proteins on the cell phenotype is dependent on binding to Cdc42, we used the CRIB (I₂₃A, S₂₄A) mutant of Borg3, which is defective in Cdc42 binding (Fig. 3B). Surprisingly, when transfected into NIH 3T3 fibroblasts, the CRIB-defective mutant induced the same dramatic phenotype as did wild-type HA₃-Borg3, with loss of stress fibers and the extension of large, protrusive lamellipodia (Fig. 7b and e). The expression of a C-terminal deletion of Borg3 (residues 1 to 83) gave a weaker phenotype, similar to the one obtained when Borg1 is overexpressed. Few cells produced protrusive lamellipodia, but many cells extended long processes (Fig. 7c). Although the CRIB-defective mutant was expressed at levels equivalent to those of the wild-type HA₃-Borg3, the Borg3(1–83) fragment did not express well, and only few cells expressed the protein at a high level. However, no degradation of the protein could be detected (Fig. 7g). The data suggest that the characteristic phenotype induced by expression of Borg3 is independent of its interaction with Cdc42.

Expression of Borg proteins delays cell spreading in a CRIB domain-dependant manner. Recent evidence has implicated several Rho family GTPases in cell motility and in spreading on fibronectin-coated surfaces (11, 34). Engagement of integrin receptors by fibronectin results in a rapid and dramatic activation of Cdc42 and Rac, which promotes their interaction with a plethora of downstream effectors (11, 55). The function of many of these interactions in motility and spreading remains

unclear. To determine whether the expression of Borg proteins would affect the ability of cells to spread, NIH 3T3 fibroblasts were transfected with either the empty pKH₃ vector, pKH₃-Borg1, or pKH₃-Borg3. Forty hours after transfection, cells were trypsinized, maintained in suspension for 15 to 20 min, and then replated onto fibronectin-coated slides. Cells were fixed with paraformaldehyde at intervals and processed for immunofluorescence labeling. Spreading was quantitated by measuring total cell surface areas of transfected or control cells. Results are shown in Fig. 8. At early times after replating (18 min), control cells were rounded, with lamellipodia encircling the cell bodies. Cells were at this stage devoid of stress fibers (Fig. 8a, untransfected cells). On the other hand, many of the cells expressing HA₃-Borg1 or HA₃-Borg3 remained very small (cell area < 500 μ m²), whereas almost no control cells were of that size (Fig. 8m). By 45 min, control cells were still rounded but had formed some stress fibers (Fig. 8b). In contrast, although the Borg-expressing cells began to spread, most of them were still small and completely devoid of stress fibers. Three hours after replating, control cells possessed abundant stress fibers and had attained a normal fibroblastic shape (Fig. 8c). Curiously, although the majority of Borg-expressing cells had by 3 h reached surface areas similar to those of the control (Fig. 8m), they exhibited an unusual phenotype, with a highly convoluted perimeter. These cells contained few stress fibers (Fig. 8c).

These results suggest that the ectopic expression of Borg1 or Borg3 can interfere with the normal process of spreading, both by delaying the rate of spreading and by interfering with the attainment of a normal fibroblastic shape after spreading is complete. The Rho GTPase is initially inhibited after engagement of integrin receptors by fibronectin but is activated at later stages of cell spreading. The effect of the Borg3 on cell shape is therefore consistent with the proposal that the Borgs partially inhibit Rho function.

As discussed above, the loss of stress fibers and changes in cell shape induced by Borg3 are independent of Cdc42 binding.

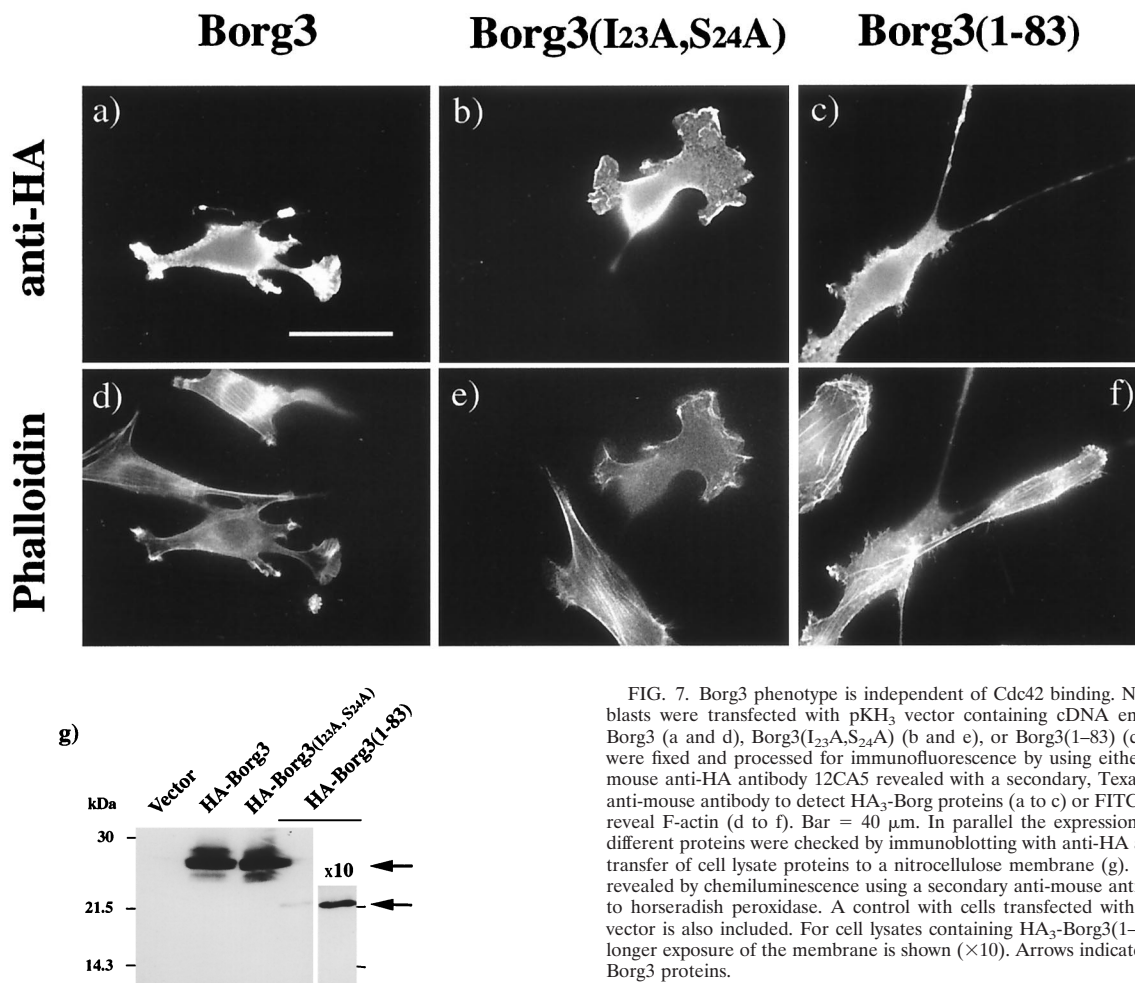


FIG. 7. Borg3 phenotype is independent of Cdc42 binding. NIH 3T3 fibroblasts were transfected with pKH₃ vector containing cDNA encoding either Borg3 (a and d), Borg3(I₂₃A,S₂₄A) (b and e), or Borg3(1-83) (c and f). Cells were fixed and processed for immunofluorescence by using either monoclonal mouse anti-HA antibody 12CA5 revealed with a secondary, Texas red-coupled anti-mouse antibody to detect HA₃-Borg proteins (a to c) or FITC-phalloidin to reveal F-actin (d to f). Bar = 40 μm. In parallel the expression levels of the different proteins were checked by immunoblotting with anti-HA antibody after transfer of cell lysate proteins to a nitrocellulose membrane (g). Proteins were revealed by chemiluminescence using a secondary anti-mouse antibody coupled to horseradish peroxidase. A control with cells transfected with pKH₃ empty vector is also included. For cell lysates containing HA₃-Borg3(1-83), a 10-fold longer exposure of the membrane is shown (×10). Arrows indicate the different Borg3 proteins.

To determine whether the reduction in spreading rate or alteration in cell shape is mediated by a Borg3-Cdc42 interaction, we assessed the effect of the mutant protein Borg3(I₂₃A,S₂₄A) and of Borg3(1-83). Ectopic expression of HA₃-Borg3(I₂₃A,S₂₄A) significantly reduced the delay in cell spreading seen with the wild-type HA₃-Borg3 (Fig. 8g and h). Within 18 min following replating, 48% of the cells expressing HA₃-Borg3(I₂₃A,S₂₄A) were larger than 10³ μm² (Fig. 8n), a proportion similar to that for the control cells (54%) and substantially higher than that for the cells expressing wild-type HA₃-Borg3 (13%). These differences were very highly significant, as determined by an unpaired *t* test (equal variances). Borg3(1-83) gave an intermediate result. The effects of the two mutants are not additive since the results obtained with a double mutant, HA₃-Borg3(I₂₃A,S₂₄A,1-83), was similar to that of cells expressing the CRIB(I₂₃A,S₂₄A) mutant (not shown). The differences in spreading rate were not due to differences in expression levels of the various proteins, as judged by quantitation of the immunofluorescence of transfected cells counted in the assay (not shown).

These data indicate that the inhibition of spreading by Borg3 depends largely on its ability to bind Cdc42 and to a lesser extent on an intact BH3 and/or Pro-rich domain. The inhibition may be partially explained by a competition with other downstream effectors for Cdc42, which is rapidly activated upon integrin receptor engagement by fibronectin.

Contrary to the effects on spreading rate, however, the two Borg3 mutants remained fully capable of interfering with the attainment of a normal fibroblastic cell shape. As can be seen in Fig. 8i and l, the transfected cells after 3 h on fibronectin possess highly convoluted, lace-like peripheries with several holes near their edges. These cells also lack stress fibers (not shown). This result is consistent with the possibility that a domain near the N terminus of Borg3 (e.g., BH1) can inhibit Rho function independently of Cdc42 binding.

Dynamic regulation of Rho is also required for cell migration (34). We therefore examined whether the ectopic expression of HA₃-Borg3 could inhibit the movement of cells into a cleared area of a monolayer. This wounding assay is expected to be less sensitive to small changes in signaling functions, because it occurs over a prolonged time period (24 h). Thus, cells that have a reduced motility rate might still have time to migrate into the cleared area. Nonetheless, as shown in Table 1, HA₃-Borg1, HA₃-Borg3, and HA₃-Borg3(I₂₃A,S₂₄A), when coexpressed with GFP, did significantly reduce the fraction of green fluorescent cells that migrated.

DISCUSSION

We have described a new family of proteins, named Borg1 to Borg5, that are putative downstream effectors of Cdc42 and/or TC10. Borg1, Borg4, and Borg5 were identified from a two-

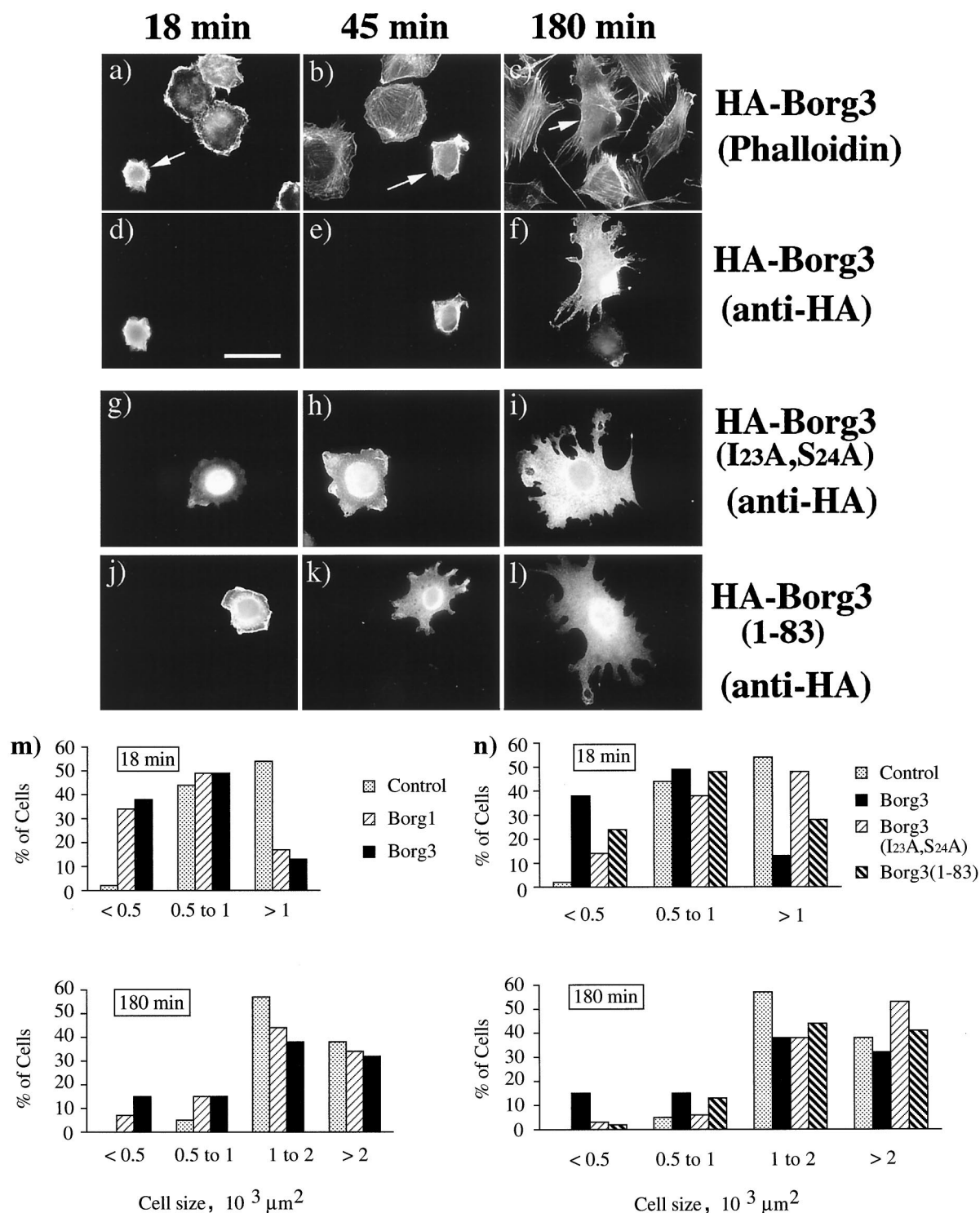


FIG. 8. Borg3 expression delays cell spreading of NIH 3T3 cells on fibronectin-coated surface in a Cdc42-binding-dependent manner. NIH 3T3 fibroblasts were transfected with pKH3 empty vector or with pKH3 containing cDNA encoding either Borg1, Borg3, Borg3(I₂₃A,S₂₄A), or Borg3(1–83); 40 h after transfection, cells were trypsinized and replated on fibronectin-coated slides. Cells were fixed and processed for immunofluorescence at specific times. FITC-phalloidin was used to reveal F-actin (a to c), and anti-HA antibody 12CA5 and a secondary, Texas Red-coupled anti-mouse antibody were used to reveal Borg proteins (d to l). Arrows in panels a to c show HA₃-Borg3-expressing cells. Bar = 40 μm . For quantitation of cell surface area at 18 and 180 min following replating (m and n), cells were visualized by FITC-phalloidin (Control) or anti-HA staining. Measurement were made on 100 cells for each time point, using the Openlab 2.0 software. Cells were then grouped according to size. Results of one representative experiment are shown. Two to four independent experiments were carried out. Data for the 18-min time point were analyzed by an unpaired *t* test. Variances were approximately equal for all data sets. Mean areas for Borg1- and Borg3-transfected cells were very highly significantly different from those of the control ($P < 0.001$) and Borg3(I₂₃A,S₂₄A)-transfected ($P < 0.001$) cells.

TABLE 1. Effects of ectopic expression of Borg proteins on fibroblasts migration in a wound test

Protein expressed with GFP	Ectopic expression (% of migrating cells expressing GFP/% of cells expressing GFP in the population)	No. of experiments
None (Vector)	1.21 \pm 0.23	2
HA-Borg1	0.77 \pm 0.13 ^a	3
HA-Borg3	0.77 \pm 0.17 ^b	3
HA-Borg3(I ₂₃ A,S ₂₄ A)	0.88 \pm 0.13 ^b	3

^a Value is significantly different from control, $P < 0.025$ (two-sample *t* test for independent samples).

^b $P < 0.05$.

hybrid screen using the TC10 GTPase as bait; Borg2 and Borg3 were discovered from database searches for Borg-related ESTs. Borg5 is most likely the human homolog of a previously reported protein called MSE55 that was shown to interact with Cdc42 and Rac through a conserved CRIB domain (3). Each member of the Borg family contains a highly related CRIB domain near the N terminus of the protein and several further regions of close similarity to one another, which we have called BH1, BH2, and BH3 domains, that were not found in any other proteins in the database. Remarkably, although Borg1, Borg2, Borg4, and Borg5 appear to bind with similar affinities to both Cdc42 and TC10, Borg3 does not interact detectably with TC10. Compared with the other members of the family, the Borg3 protein sequence lacks three amino acids immediately upstream of the CRIB domain. This deletion might account for the differential binding to the small GTPases. This result extends our previous observations that closely related pairs of effectors, such as MLK2-MLK3 and WASP-N-WASP, interact differentially with TC10 (33). TC10 has also been reported to be insensitive to the nucleotide exchange factor Dbl (17). Thus, TC10 may be designed to respond in a restricted way to only a subset of upstream signals and to activate only a specific subset of the downstream effectors that are regulated by Cdc42. Such restrictions may be essential if TC10 operates mainly in highly differentiated tissues such as muscle, where proliferative signals could be deleterious to function.

The tissue distribution of Borgs varies with the isoform. Borg1 mRNA was detected in all tissues examined, while Borg2 was expressed only at very low levels in brain and liver. Borg3 mRNA appears to be present at very low levels in most tissues, but a much longer transcript was clearly detectable in skeletal muscle. Whether the multiple Borg transcripts represent splice variants or partially processed mRNAs remains to be determined. We were unable to clone the full-length sequence of Borg4, and no mRNA was detected for this protein in any of the adult human tissues tested. It may therefore be expressed only in fetal or embryonic tissues.

We studied Borg1 and Borg3 in more detail. When expressed ectopically in NIH 3T3 cells (or in BHK cells), the proteins appeared to be predominantly cytoplasmic and induced a characteristic phenotype that included the formation of long thin extensions and protrusive lamellipodia as well as the partial loss of stress fibers. The effect may be related to the loss of stress fibers caused by inhibition of Rho function, which permits a more extensive remodeling of the F-actin into filopodia (35). The idea that there is competition between different GTPase-mediated signals for structuring the finite amount of actin present within the cell has been also proposed by others (32). We tested this hypothesis further by coexpressing Borg3

with wild-type Rho. The overexpression of Rho restored stress fibers and a more normal fibroblastic cell shape, supporting the idea that Borg3 can inhibit Rho function.

By what mechanism might the Borg proteins exert their effects on the cell? Conceivably, the Borg CRIB domain might bind to Cdc42 and/or TC10 and compete out other downstream effectors for these GTPases. We do not favor this mechanism, however, because a CRIB mutant of Borg3 that is unable to bind Cdc42 was fully capable of inducing the characteristic Borg phenotype. The CRIB mutant data also rule out the possibility that the phenotype might result from the interaction of GTP-bound Cdc42 and/or TC10 with Borg, which activates a downstream signal. Moreover, the coexpression of Borg3 with wild-type Cdc42 did not inhibit the formation of filopodia by Cdc42 but caused a dramatic increase in their length. A similar phenotype could be induced by the coexpression of the RhoGAP, p190, with Cdc42, supporting the proposal that Borgs can inhibit Rho function. Therefore, the Borg proteins most likely induce the observed effects on the cytoskeleton independently of a direct interaction with the small GTPases, although binding to Cdc42 may well modify these effects, either by changing the subcellular distribution of the Borg proteins or by inducing conformational changes which would alter their interactions with other proteins.

It is important to note that changes in stress fiber formation can occur independently of Rho activity. For example, the drug cytochalasin D activates Rho but causes the dissolution of stress fibers (38). Rho is believed to regulate stress fiber formation through its downstream effector, ROCK, which phosphorylates and inhibits the myosin light-chain (MLC) phosphatase. Phosphorylation of MLC mediates the formation of actomyosin bundles and stimulates contraction (21). A protein kinase effector of Cdc42 and Rac (PAK1) has been reported to phosphorylate and inhibit the MLC kinase, thereby reducing MLC phosphorylation and reducing actomyosin bundling (45), but these results have been disputed, and in another system PAK1 appears to be able to stimulate MLC phosphorylation (47). Our present data do not, therefore, definitively prove that Borg proteins cause disassembly of stress fibers through an inhibition of Rho function, but they are consistent with this hypothesis.

The cytoskeletal responses of cells to Rac have been shown to be independent of the effects of this GTPase on other signaling pathways such as that which activates JNK (24). A similar situation probably holds for other Rho family GTPases, including Cdc42. To determine whether Borg proteins participate in these other pathways, we examined whether the expression of Borg3 interferes with or potentiates JNK activation. We showed that wild-type HA₃-tagged Borg3 inhibited the basal JNK activity. Surprisingly, like its effects on the cytoskeleton, the Borg3-mediated inhibition of JNK was independent of its ability to bind Cdc42. It remains to be determined whether the mechanism by which the Borg protein interferes with JNK is the same as that by which it inhibits Rho function.

Our observations on the effects of Borg3 on cell spreading revealed two distinct responses, only one of which was independent of Cdc42 binding. After replating onto fibronectin, the cells that expressed HA₃-Borg3 spread very slowly, and the spread cells possessed an unusual cell shape and lacked stress fibers. Using the CRIB mutant which is defective in Cdc42 binding, we were able to partially restore the normal spreading rate but did not restore normal cell shape. We propose that the rate of spreading is at least partially dependent on the activation of Cdc42 and its interaction with a set of downstream effectors, which may be inhibited by competition with the Borg

protein. WASP and N-WASP interact with the Arp2-Arp3 complex. This complex, which enhances actin nucleation, branching, and cross-linking of actin filaments *in vitro*, is involved in the formation of lamellipodia (27, 42). However, competition cannot completely account for the inhibition in cell spreading, because expression of the mutant Borg3(1–83), which can bind Cdc42, was also relatively ineffective in inhibiting spreading. Borg3(1–83) is expressed much less efficiently than wild-type protein, but even when only cells that expressed similar amounts of protein were compared, the Borg3(1–83) mutant did not show the same degree of inhibition of spreading as did the wild-type protein. It is likely, therefore, that protein-protein interactions involving the C-terminal domains of Borg are required, in addition to Cdc42 binding, to inhibit cell spreading on fibronectin. The abnormal cell shape caused by expression of either the CRIB mutant or the C-terminal deletion mutant is consistent with our other data that implicate Borgs in Rho inhibition and with evidence from other groups that Rho is switched on at a late stage in cell spreading and may be necessary for the assumption of a normal fibroblastic cell shape (5, 11).

The wounding assay for cell motility was less sensitive to ectopic expression of Borg3. Although a small inhibition was consistently observed, it was much less dramatic than the inhibition of spreading. This difference may reflect the very different times over which the two experiments were conducted. Motility was estimated after 24 h, during which time the cell could compensate for partial inhibition of endogenous functions either by changes in gene expression or by adaptation in the responsiveness to external signals.

Overall, our studies suggest that Borg proteins may interfere with the activity of the Rho GTPase and function as negative regulators in cross talk between Cdc42- and Rho-mediated signaling pathways. The mechanism by which Borg proteins inhibit actin stress fiber formation remains to be determined. They could inhibit signaling through the lysophosphatidic acid receptor or through the G α proteins that couple to Rho-specific guanine nucleotide exchange factors, or they could inhibit the exchange factors themselves. Alternatively, they may interfere with the phosphorylation of MLC. It will be of interest, therefore, to identify binding partners for the Borg proteins other than Cdc42 and TC10 as a first step toward a complete description of their function.

ACKNOWLEDGMENTS

We thank Stan Hollenberg, Rick Cerione, Mark Rush, and Gary Bokoch for their kind gifts of reagents, and we thank Cheryl Neudauer, Nia Tatsis, and Tom Parsons for helpful discussions.

This work was supported by the Markey Center for Cell Signaling and by grant CA 56300 from the National Institutes of Health.

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